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AMENDMENTS TO THE CLAIMS

This listing of claims replaces all prior listing of claims in the application.

1. (Presently amended) A compound of Formula I or Formula II

or pharmaceutically acceptable salt thereof, wherein

R1 is selected from the group consisting of

- (a) phenyl, optionally substituted at positions 3 and 4 halogens,
- (b) -O-isopropyl,
- (c) -O-cyclopropyl, and
- (d) -O-CH2-cyclopropyl;

R² is selected from the group consisting of:

- (a) -S(O)2CH3, and
- (b) -S(O)2NH2;

 $\ensuremath{\mathrm{R}}^3$ is selected from the group consisting of

- (a) hydrogen,
- (b) methyl,
- (c) ethyl,
- (d) hydroxyl,

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- (e) F, Cl, and
- (f) CF3;

R4 is selected from the group consisting of

- (a) methyl, and
- (b) ethyl;

X1 is selected from the group consisting of:

- (a) -OCH2-,
- (b) $-OC(R^3)(R^4)$ -,
- (c) -CH2-linker -O-, and
- (d) $-C(R^3)(R^4)$ -linker-O-,

wherein the oxygen end of X1 is attached to the carbonyl carbon of Formula I;

X2 is selected from the group consisting of:

- (a) --OCH2--,
- (b) $-OC(R^3)(R^4)$ -,
- (c) -CH2-linker -O-, and
- (d) -C(R³)(R⁴)-linker-O-;

wherein the carbon end of X^2 is attached to the carbon adjacent to the R^2 -phenyl explicitly shown:

- -linker is selected from the group consisting of
 - (a) -C(O)-(CH2)m-O-,
 - (b) -C(O)-(CH2)m(-O-(CH2)n)p-O-, and
 - (c) -C(O)-aryl-O-,
 - (d) C(O) heteroaryl O.

wherein m, n and p are each independently integers ranging from 0 to 6;

Y is selected from the group consisting of

- (a) hydrogen, and
- (b) acyl,

wherein the acyl group is selected from the group consisting of

- (a) -C(O) -C₁-6alkyl, optionally substituted with 1, 2 or 3 substituents independently selected from the group consisting of halo, hydroxyl, amino, C₁-3alkoxy, aminoC₁-3alkyl,
- (b) −C(O) −aryl, <u>and</u>

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(d) (c) an amino acid;

Z is selected from the group consisting of:

- (a) -OR⁵,
- (b) -NR5R6,

wherein R5 and R6 are each independently selected from

- (a) hydrogen,
- (b) C₁₋₆alkyl,
- (c) phenyl, and(d) C₁₋₂-phenyl,

wherein R⁵ and R⁶ choices (b), (c) and (d) are optionally substituted with 1, 2, or 3 substituents selected from halo, hydroxyl, amino, C₁₋₃alkyl, and G₁₋₃alkoxy; C₁₋₃alkoxy;

X is selected from the group consisting of:

- (a) -OCH2-, and
- (b) $-C(R^3)(R^4)O-$,

wherein the carbon at the end of X is attached to the carbon adjacent to the phenyl;

 Y^1 is -linker1-, which is selected from the group consisting of

- (a) -C(O)-(CH₂)_r-C(O)-,
- (b) -C(O)-aryl-C(O)-,
- (c) C(O) heteroaryl C(O),
- (d) (c) -C(O)-(CH₂)_r-(O-(CH₂)_s)_t-C(O)-, and
- (e) (d) -C(O)-(CH₂)_r-CH-(CH₂)_s-C(O)-,

wherein r, s and t are each independently integers ranging from 0 to 6. $\underline{6}$; and

Z1 is selected from the group consisting of:

- (a) -OR⁵, <u>and</u>
- (b) -NR5R6.
- (Original) A compound according to claim 1 of Formula I

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- 3. (Original) A compound according to claim 2 wherein: \mathbb{R}^1 is phenyl, optionally substituted at positions 3 and 4 with fluorine.
- 4. (Original) A compound according to claim 2 wherein: R^2 is $-S(O)_2CH_3$.
- $\label{eq:continuous} 5, \qquad \text{(Original) A compound according to claim 2 wherein:} \\ R^3 \text{ is selected from the group consisting of}$
 - (a) hydrogen,
 - (b) methyl, and
 - (c) ethyl.
- $6. \qquad \hbox{(Original) A compound according to claim 2 wherein:} X^1 and X^2 are each is selected from the group consisting of:}$
 - (a) -OCH2-, and
 - (b) $-OC(R^3)(R^4)$ -.
- 7. (Original) A compound according to claim 2 wherein: Y is hydrogen or -OCH3.
- 8. (Original) A compound according to claim 2 wherein: Z is hydroxyl or -OCH3.
- 9. (Original) A compound according to claim 2 wherein: R^1 is phenyl, optionally substituted at positions 3 and 4 with fluorine; R^2 is $-S(O)_2CH_3;\\$
- \mathbb{R}^3 is selected from the group consisting of

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(a) hydrogen,

(b) methyl, and

(c) ethyl;

R4 is selected from the group consisting of

(a) methyl, and

(b) ethyl;

X1 and X2 are each is selected from the group consisting of:

(a) -OCH2-, and

(b) $-OC(R^3)(R^4)$ -;

Y is hydrogen or -OCH3; and

Z is hydroxyl or -OCH3.

10. (Original) A compound according to claim 1 of Formula II

- $11. \qquad \hbox{(Original) A compound according to claim 10 wherein:} \\ R^1 \hbox{ is phenyl, optionally substituted at positions 3 and 4 halogens.}$
- 12. (Original) A compound according to claim 11 wherein: $R^2 \ {\rm is} \ {\rm -S(O)_2CH_3}.$

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13. A compound according to claim 12 wherein:

R³ is selected from the group consisting of

- (a) hydrogen,
- (b) methyl, and
- (c) ethyl.
- 14. (Original) A compound according to claim 13 wherein: Y 1 is selected from -(O)C(H)=C(H)C(O)- and -(O)C(CH2)2C(O)-.
- 15. (Original) A compound according to claim 14 wherein: Z^1 is hydroxyl or –OCH3.
- $16. \qquad (Original)\ A\ compound\ according\ to\ claim\ 15\ wherein:$ $R^1\ is\ phenyl,\ optionally\ substituted\ at\ positions\ 3\ and\ 4\ halogens;$ $R^2\ is\ -S(O)_2CH_3;$

R³ is selected from the group consisting of

- (a) hydrogen,
- (b) methyl, and
- (c) ethyl;

 Y^1 is selected from -(O)C(H)=C(H)C(O)- and -(O)C(CH₂)₂C(O)-; and Z^1 is hydroxyl or -OCH₃.

- 17. (Cancelled)
- 18. (Cancelled)
- (Original) A pharmaceutical composition comprising a compound according to Claim 1 and a pharmaceutically acceptable carrier.
 - 20. (Presently amended) A compound according to claim 1 selected from

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